

## CLAIM AMENDMENTS

1 through 9 (canceled).

- 1           10. (New) A compound selected from the group consisting  
2 of  
3 (a) Leu Lys Ala Thr Thr Asn Ser Lys Leu Met Met Tyr (Seq ID NO: 1);  
4 (b) Val Asp Met Ile Asn Asp Val Gln Pro Leu Thr Pro (Seq ID NO: 2);  
5 (c) Val Asp Met Ile Asp Asp Val Gln Pro Leu Thr Pro (Seq ID NO: 3);  
6 (d) Val Asp Met Ile Asn Asp Val Gln Pro Met Thr Pro (Seq ID NO: 4);  
7 (e) Val Tyr Met Met Asn Asn Gly Gln Pro Pro Ser Pro (Seq ID NO: 5);  
8 (f) Val Asp Met Ile Asn Asp Val Gln Pro Met Ser Pro (Seq ID NO: 6);  
9 (g) Trp His Trp Gln Trp Thr Pro Trp Ser Ile Gln Pro (Seq ID NO: 7);  
10 (h) His Ser Pro Leu Asp Ser Ser Arg His Ala Thr Tyr (Seq ID NO: 8);  
11 (i) His Tyr Thr Leu Asp Ser Cys Arg His Pro Thr Tyr (Seq ID NO: 9);  
12 (j) Val Tyr Ser Ser Thr Thr Arg Pro Leu Pro Ser Pro (Seq ID NO: 10);  
13 (k) Val Tyr Ser Ser Asn Thr Arg Pro Leu Pro Ser Pro (Seq ID NO: 11);  
14 (l) Val Tyr Ser Ser Asn Asn Arg Pro Leu Pro Ser Pro (Seq ID NO: 12);  
15 (m) Val Tyr Leu Leu Asn Asn Arg Pro Leu Pro Ser Pro (Seq ID NO: 13);  
16 (n) Val Tyr Leu Leu Ser Thr Arg Pro Leu Pro Ser Pro (Seq ID NO: 14);  
17 (o) Val Tyr Trp Pro Thr Asn Arg Pro Leu Pro Ser Pro (Seq ID NO: 15);  
18 (p) Val Gln Pro Ser Ile Asn Arn Pro His Gln Arg Pro (Seq ID NO: 16);  
19 (q) Tyr His Asn Tyr Thr Thr Ala Pro His Ser Pro Ser (Seq ID NO: 17);  
20 (r) Lys Pro Val Ile Ser Pro Thr Asn Ala Leu Gln Pro (Seq ID NO: 18);  
21 (s) Val Thr Gly Pro Thr Lys Asn Leu Pro Ala Thr Thr (Seq ID NO: 19);  
22 (t) Ala Ser His Val Asp Tyr Arg Arg Phe Leu Leu Thr (Seq ID NO: 20);  
23 (u) Asp Gln Asp Phe Als Pro Asp Arg His Tyr Arg Leu (Seq ID NO: 21);  
24 (v) Gln Lys Trp Pro Glu Thr Tyr Pro Asp Leu Ser Phe (Seq ID NO: 22);

25 (w) Gly Asp Pro Val Pro Gln Thr Tyr Ser Ala Ala Gly (Seq ID NO: 23);  
26 (x) Ala Val Ser Val Asn Thr Lys Ile Asp Thr Glu Ala (Seq ID NO: 24);  
27 (y) Gln Pro Asn Tyr Thr Ser Leu Leu Tyr Gly Thr Glu (Seq ID NO: 25);  
28 (z) Thr Gln Pro Pro Ile His His Tyr Gln Leu Pro Ala (Seq ID NO: 26);  
29 and  
30 (aa) Gly Trp Asp His Ile His Gly Val His Gln His Val (Seq ID NO:  
31 27)).

1 11. (New) Leu Lys Ala Thr Thr Asn Ser Lys Leu Met Met Tyr  
2 (Seq ID NO: 1) as defined in claim 10.

1 12. (New) Val Asp Met Ile Asn Asp Val Gln Pro Leu Thr Pro  
2 (Seq ID NO: 2) as defined in claim 10.

1 13. (New) His Ser Pro Leu Asp Ser Ser Arg His Ala Thr Tyr  
2 (Seq ID NO: 8) as defined in claim 10.

1 14. (New) Val Tyr Ser Ser Thr Thr Arg Pro Leu Pro Ser Pro  
2 (Seq ID NO: 10) as defined in claim 10.

1 15. (New) A pharmaceutical composition for the treatment  
2 of transmissible spongiform encephalopathy which comprises a  
3 therapeutically effective amount of the compound defined in claim 1  
4 together with a pharmaceutically acceptable inert carrier or  
5 diluent.

1           16. (New) The pharmaceutical composition defined in claim  
2   15 in solid, semiliquid or liquid form.

1           17. (New) The pharmaceutical composition defined in claim  
2   15 in the form of an injection solution, drop, juice, syrup, spray,  
3   suspension, granulate, tablet, pellet, transdermal therapeutic  
4   system, capsule, plaster, suppository, salve, cream, lotion, gel,  
5   emulsion or aerosol form.

1           18. (New) The pharmaceutical composition defined in claim  
2   15 further comprising an auxiliary substance selected from the group  
3   consisting of a surface active substance, a coloring agent, a  
4   preservative, a bursting agent, a smoothing agent, a lubricant, an  
5   aromatizing agent and/or a binder.

1           19. (New) The peptide defined in claim 10 substituted or  
2   modified by at least one component selected from the group  
3   consisting of sugar residues, glucoromic acid, sulfate residues,  
4   serine, glycine or aspartate.

1           20. (New) A method of making a peptide as defined in  
2   claim 10 in that a solid phase synthesis in a liquid phase is used.

1           21. (New) A method of making a peptide as defined in  
2   claim 10 wherein the peptide is expressed by a nucleic acid coding  
3   therefore.

1           22. (New) A method of inhibiting replication of a PrP<sup>Sc</sup>  
2     prion in a mammalian subject which comprises the step of  
3     administering to said subject a therapeutically effective amount of  
4     the compound defined in claim 10.

1           23. (New) The method of inhibiting replication of a PrP<sup>Sc</sup>  
2     prion in a mammalian subject defined in claim 22 wherein the  
3     mammalian subject is a human.